## **The Claims**

This listing of claims will replace all prior versions and listings of claims in the application:

- 1. (Currently amended) A process for hydrogen peroxide plasma sterilization comprising:
- (a) inserting at least one primary container containing a temperature-sensitive material into a sterilization treatment chamber;
  - (b) lowering the pressure in the treatment chamber to create a vacuum;
  - (c) injecting, at least one time, hydrogen peroxide into the chamber;
  - (d) lowering the pressure in the treatment chamber to reestablish a vacuum;
  - (e) generating a plasma; and
  - (f) ventilating the chamber;

wherein the chamber temperature is less than 39°C throughout the process, the temperature-sensitive material is sterilized and stable, and wherein the pressure in step (d) is about 100 to 800 mtorr.

- 2. (Previously presented) The process of claim 1, wherein the pressure in step (b) is about 100 to 800 mtorr.
- 3. (Previously presented) The process of claim 1, wherein step (c) is performed from between 1 and 60 minutes.
- 4. (Previously presented) The process of claim 1, wherein prior to step (d) a hydrogen peroxide diffusion step is performed simultaneously with ventilation.
- 5. (Previously presented) The process of claim 1, wherein prior to step (d) a hydrogen peroxide diffusion step is performed without ventilation.

- 6. (Previously presented) The process of claim 4, wherein the hydrogen peroxide diffusion step is performed from between 1 and 60 minutes.
- 7. (Previously presented) The process of claim 5, wherein the hydrogen peroxide diffusion step is performed from between 1 and 60 minutes.
- 8. (Previously presented) The process of claim 1, wherein the temperature of the temperature-sensitive material does not rise above 40°C during the sterilization process.
- 9. (Previously presented) The process of claim 1, wherein the temperature-sensitive material comprises biological materials.
- 10. (Previously presented) The process of claim 9, wherein the biological materials are proteins, peptides, nucleic acids, lipids, or cellular materials.
- 11. (Previously presented) The process of claim 9, wherein the biological material is a fibrogen containing solution.
- 12. (Previously presented) The process of claim 9, wherein the biological material is a Factor XIII containing solution.
- 13. (Previously presented) The process of claim 9, wherein the biological material is a thrombin containing solution.
- 14. (Previously presented) The process of claim 9, wherein the biological material comprises the components of tissue glue.
- 15. (Previously presented) The process of claim 9, wherein the biological material comprises the components of fibrin glue.
- 16. (Previously presented) The process of claim 1, wherein the temperaturesensitive material comprises non-biological materials.

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- 17. (Previously presented) The process of claim 1, wherein the process is performed more than one time.
- 18. (Previously presented) The process of claim 1, wherein before step (b), a preplasma step is performed comprising:
  - lowering the pressure in the treatment chamber to create a vacuum;
  - applying a preplasma; and
  - ventilating the chamber.
- 19. (Previously presented) The process of claim 18, wherein the pressure is about 100 to 800 mtorr.
- 20. (Previously presented) The process of claim 18, wherein the preplasma is applied for about 1 to 30 minutes.
- 21. (Previously presented) The process of claim 18, wherein the ventilation step is no greater than 5 minutes.
- 22. (Previously presented) The process of claim 1, wherein the primary container is enveloped at least one time with materials partially permeable to hydrogen peroxide.
- 23. (Previously presented) The process of claim 1, wherein the primary container containing the temperature-sensitive material is placed in a secondary container.
- 24. (New) The process of claim 1, wherein the chamber temperature is between 20-39°C.
- 25. (New) The process of claim 24, wherein the chamber temperature is between 25-35°C.